

MAR 30 2005

510(k) Summary

Submitted By:

Stephanie Roberts
Regulatory Affairs
Cook Incorporated
750 Daniels Way, PO Box 489
Bloomington, IN 47402
812-339-2235

March 17, 2005

Device:

Trade Name: Zilver® 518™ Biliary Stent
Proposed Classification: Catheter, Biliary, Diagnostic

Predicate Devices:

The Zilver® 518™ Biliary Stent is similar in terms of intended use, materials of construction and technological characteristics to predicate devices reviewed as devices for palliation of malignant neoplasm in the biliary tree.

Device Description:

The Zilver® 518™ Biliary Stent is a self-expanding, nitinol stent designed for excellent radial strength and optimal longitudinal flexibility. Constructed from a series of interconnected Z-shaped segments, the stent conforms to the shape of the biliary system and provides circumferential scaffolding throughout the stent's length. Gold radiopaque markers on each end of the Zilver® 518™ Biliary Stent, along with radiopaque markers on the delivery system, allow precise positioning of the stent. The stent's interconnected Z-shaped segments also keep foreshortening to a minimum.

The Zilver® 518™ Biliary Stent comes preloaded in a 5 French delivery system. The stent is deployed with the use of a simple hand held device. The stent is currently available in unrestrained outer diameters of 4, 5, 6, 7, 8, 9 and 10 mm in lengths of 20, 30, 40, 50 and 60 mm. The 80 mm length stent is an addition to this product line.

Substantial Equivalence:

This device will be manufactured according to specified process controls and a Quality Assurance Program. This device will undergo packaging similar to the predicate devices currently marketed and distributed by Cook Incorporated. This device will undergo sterilization similar to the predicate devices currently marketed and distributed. Being similar with respect to indications for use, materials and physical construction to predicate devices, this device meets the requirements for section 510(k) substantial equivalence.

Test Data:

The Zilver[®] 518™ Biliary Stent was subjected to the following tests to assure reliable design and performance under the specified testing parameters. These tests were comprised of:

1. Deployment
 - a. Profile
 - b. Deployment Accuracy
 - c. Stent length and Change due to Deployment
 - d. Stent Diameter
 - e. Uniformity of expansion
 - f. Stent Integrity

The results of these tests provide reasonable assurance that the device has been designed and tested to assure conformance to the requirements for its use as a biliary stent.



DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAR 30 2005

Ms. Stephanie Roberts
Regulatory Affairs Specialist
Cook® Incorporated
P.O. Box 489
BLOOMINGTON IN 47402-0489

Re: K050698
Trade/Device Name: Zilver® 518™ Biliary Stent
Regulation Number: 21 CFR §876.5010
Regulation Name: Biliary catheter and accessories
Regulatory Class: II
Product Code: FGE
Dated: March 17, 2005
Received: March 17, 2005

Dear Ms. Roberts:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act and the limitations described below. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

The Office of Device Evaluation has determined that there is a reasonable likelihood that this device will be used for an intended use not identified in the proposed labeling and that such use could cause harm. Therefore, in accordance with Section 513(i)(1)(E) of the Act, the following limitation must appear in the Warnings section of the device's labeling:

The safety and effectiveness of this device for use in the vascular system have not been established.

Furthermore, the indication for biliary use must be prominently displayed in all labeling, including pouch, box, and carton labels, instructions for use, and other promotional materials, in close proximity to the trade name, of a similar point size, and in bold print.

Page 2 – Ms. Stephanie Roberts

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and permits your device to proceed to the market. This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification if the limitation statement described above is added to your labeling.

Please note that the above labeling limitations are required by Section 513(i)(1)(E) of the Act. Therefore, a new 510(k) is required before these limitations are modified in any way or removed from the device's labeling.

If you desire specific information about the application of other labeling requirements to your device (21 CFR Part 801), please contact the Office of Compliance at (301) 594-4616. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International, and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>.

Sincerely yours,

A handwritten signature in black ink, appearing to read "Donna-Bea Tillman".

Donna-Bea Tillman, Ph.D.
Director
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

510(k) Number: K050698

Device Name: Zilver[®] 518[™] Biliary Stent

FDA's Statement of the Indications for Use for device:

The Zilver[®] 518[™] Biliary Stent is indicated for use in the palliation of malignant strictures (neoplasms) in the biliary tree.

Prescription Use ✓
(Per 21 CFR 801.109)

OR

Over-the-Counter Use _____

Jane C. Bragdon
(Division Sign-Off)
Division Reproductive, Abdominal,
and Radiological Devices
510(k) Number K050698